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July 13, 1999



Office of Special Nutritionals (HFS-450) Center for Food Safety and Applied Nutrition Food and Drug Administration 200 C Street, SW Washington, DC 20204

RE: <u>Notification of Additional Dietary Supplement Marketing Claims</u>

Dear Sir/Madam:

Pursuant to 21 CFR § 101.93 this is to notify you that DCV, Inc. and its wholly owned subsidiary Legacy USA, Inc. will be marketing a dietary supplement containing the structure/function claims as indicated below.

1. Name and Address of Manufacturer and Distributor:

Manufacturer:
Phoenix Labs
140 Lauman Avenue
Hicksville, NY 11801

<u>Distributor:</u> Legacy **USA**, Inc 102 Harbor City Blvd. Melborne, FL 32901

2. <u>Text of Statement:</u>

The following statements will be made in marketing the product:

"This product:

- a) helps inhibit cartilage deterioration,
- b) promotes joint lubrication,
- c) helps maintain joint mobility, and
- d) supports cartilage growth."

3. <u>Description of Dietary Ingredient or Supplement:</u>

The product is a capsule containing glucosamine hydrochloride.

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4. <u>Name of **Dietary** Supplement:</u>

BioChoice® FlexTM

5. <u>Certification of Accuracy:</u>

I, Neal Kane, Vice President of DCV, Inc. do hereby **certify** that the information contained in this notice is complete and accurate and that DCV, Inc. has **substantiation** that the statement to be made is **truthful** and not misleading.

Neal J. Kane